Claims.

1. A 6-mercapto-cyclodextrin derivative having the general formula I

Formula I

wherein m is 0-7 and n is 1-8 and m+n = 7 or 8;

R is (C_{1-6}) alkylene, optionally substituted with 1-3 OH groups,

or (CH₂)_o-phenylene-(CH₂)_p;

o and p are independently 0-4;

X is COOH, CONHR₁, NHCOR₂, SO₂OH, PO(OH)₂, O(CH₂-CH₂-O)_q-H,

OH or tetrazol-5-yl;

R₁ is H or (C₁₋₃)alkyl;

R₂ is carboxyphenyl;

q is 1-3;

or pharmaceutically acceptable salts thereof;

with the exclusion of

6-per-deoxy-6-per-(2-hydroxyethylthio)-β-cyclodextrin;

6-mono-deoxy-6-mono-(2-hydroxyethylthio)-β-cyclodextrin;

6-per-deoxy-6-per-(2-hydroxyethylthio)-γ-cyclodextrin;

6-per-deoxy-6-per-(carboxymethylthio)- β -cyclodextrin;

6-mono-deoxy-6-mono-(carboxymethylthio)-β-cyclodextrin;

 $6A,6B-dideoxy-6A,6B-bis[(o-carboxyphenyl)thio]-\beta-cyclodextrin;$

6A,6B-dideoxy-6A,6B-bis(carboxymethylthiol)-β-cyclodextrin and

6-per-deoxy-6-per-(2,3-dihydroxypropylthio)- β -cyclodextrin.

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2. The 6-mercapto-cyclodextrin derivative according to claim 1, wherein R, m and n are defined as in claim 1 and X is COOH or SO₂OH; or a pharmaceutically acceptable salt thereof.

- 3. The 6-mercapto-cyclodextrin derivative according to claim 1, wherein m is 0; n is 8; R is (C₁₋₆)alkylene or (CH₂)₀-phenylene-(CH₂)_p, o and p are independently 0-4; and X is COOH or SO₂OH; or a pharmaceutically acceptable salt thereof.
- 4. A 6-mercapto-cyclodextrin derivative according to any of claims 1-3 selected from:

6-per-deoxy-6-per-(2-carboxyethyl)thio-γ-cyclodextrin;

6-per-deoxy-6-per-(3-carboxypropyl)thio-γ-cyclodextrin;

6-per-deoxy-6-per-(4-carboxyphenyl)thio-γ-cyclodextrin;

6-per-deoxy-6-per-(4-carboxyphenylmethyl)thio-γ-cyclodextrin;

6-per-deoxy-6-per-(2-carboxypropyl)thio-γ-cyclodextrin; and

6-per-deoxy-6-per-(2-sulfoethyl)thio-γ-cyclodextrin;

or a pharmaceutically acceptable salt thereof.

- 5. A 6-mercapto-cyclodextrin derivative according to the general formula I of claim 1 for use in therapy.
- The use of a 6-mercapto-cyclodextrin derivative according to the general formula I of claim 1 for the manufacture of a medicament for the reversal of drug-induced neuromuscular block.
- 7. A kit for providing neuromuscular block and its reversal comprising (a) a neuromuscular blocking agent, and (b) a 6-mercapto-cyclodextrin derivative according to the general formula I of claim 1.
- 8. The kit according to claim 6, wherein the neuromuscular blocking agent is selected from the group consisting of rocuronium, vecuronium,

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pancuronium, rapacuronium, mivacurium, (cis)atracurium, tubocurarine and suxamethonium.

- 9. The kit according to claim 7, wherein the neuromuscular blocking agent is rocuronium.
- 10. A pharmaceutical composition comprising a 6-mercapto-cyclodextrin derivative having the general formula I

Formula I

wherein m is 0-7 and n is 1-8 and m+n = 7 or 8;

R is (C_{1.6})alkylene, optionally substituted with 1-3 OH groups,

or $(CH_2)_o$ -phenylene- $(CH_2)_p$;

o and p are independently 0-4;

X is COOH, CONHR₁, NHCOR₂, SO₂OH, PO(OH)₂, O(CH₂-CH₂-O)₀-H,

OH or tetrazol-5-yl;

R₁ is H or (C₁₋₃)alkyl;

R₂ is carboxyphenyl;

q is 1-3;

or a pharmaceutically acceptable salt thereof, in admixture with pharmaceutically acceptable auxilliaries.

11. A method for reversal of drug-induced neuromuscular block in a patient, which comprises parenterally administering to said patient an effective amount of a 6-mercapto-cyclodextrin derivative according to the general formula I of claim 1.